PREPARATION OF SUBSTITUTED CYCLOPENTANE AND

CYCLOPENTENE COMPOUNDS AND CERTAIN INTERMEDIATES

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Technical Field

This invention relates to methods for preparing certain substituted cyclopentane compounds and certain intermediates thereof. The present invention is also concerned with novel intermediates or precursors for producing the substituted cyclopentane compounds. Substituted cyclopentane compounds prepared according to the present invention are useful as neuraminidase inhibitors, and especially in pharmaceutical composition for preventing, treating or ameliorating viral, bacterial and other infections.

Background of the Invention

Despite the wealth of information available, influenza remains a potentially devastating disease of man, lower mammals, and birds. No effective vaccine exists and no cure is available once the infection has been initiated.

25 Influenza viruses consist of eight pieces of single stranded RNA, packaged in orderly fashion within the virion. Each piece codes for one of the major viral proteins. The replication complex is enclosed with a membrane composed of matrix protein associated with a lipid bilayer. Embedded in

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converting the alcohol group of said acylated compound into a leaving group;

displacing said leaving group with ammonia or guanidine to obtain said compound of formula la; or displacing said leaving group with an azide ion and then converting to a guanidine with a NH₂ compound to obtain said compound of formula la.

-4. A method for preparing isoxazoline compounds

represented by the formula 10:

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wherein each R₁ individually is alkyl or substituted alkyl, alkenyl or substituted alkenyl of 1-6 carbon atoms, or H; 5 each of R₂ and R₃ individually is alkyl or alkenyl of 1-8 carbon atoms, cycloalkyl or substituted cycloalkyl of 4-8 carbon atoms, aryl or substituted aryl, arylalkyl or substituted arylalkyl, or H provided at least one of R₂ and R₃ is other than H;

which comprises reacting a nitrite oxide of formula 2

with a cyclopentane derivative of the formula

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to produce said isoxazoline compound.

A method for preparing a substituted cyclopentane compound represented by formulae 1a or 1b

wherein each R1 individually is alkyl or substituted alkyl, alkenyl or substituted alkenyl of 1-6 carbon atoms, or H; each of R2 and R3 individually is alkyl or alkenyl of 1-8 carbon atoms, cycloalkyl or substituted cycloalkyl of 4-8 carbon atoms, anyl or substituted anyl, anylalkyl or substituted arylalkyl, or H provided at least one of R2 and R3 is other than H; X is NHR1, NHC(=NH)NHR4 where R4 is H, alkyl of 1-6 carbon atoms, OR1, COR1, COOR1 CN or NO2; A is H, F, OR1, OCOR1, -OOCNHR1, NHR1, or NHCOOR1; and pharmaceutically acceptable salts thereof; > To control

which comprises:___

obtaining an isoxazoline compound of formula according to claim 47

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compound of formula 13

converting said isoxazoline to a compound of formula 12

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and dehydrating said compound of formula 12 to produce a

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or converting the OH groups of said compound of formula 12

6. An isoxazoline derivative represented by the following formula 4:

Intermediate

wherein each of R, and R, individually is alkyl or alkenyl of 1-8 carbon atoms, cycloalkyl or substituted cycloalkyl of 4-8 carbon atoms, arylalkyl or substituted arylalkyl, or H provided at least one of R, and R, is other than H; each of Y and Z individually is COOR, or H provided that at least one of Y and Z is other than H.

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